Application No.: 10/576,693 Docket No.: 0230-0238PUS1

Amendment dated September 15, 2010 Reply to Office Action of March 16, 2010

AMENDMENTS TO THE CLAIMS

 (Currently Amended) A method for screening genomic DNA fragments capable of providing plants with an agriculturally advantageous phenotypic variation, comprising the steps of:

1) preparing genomic DNA from a <u>donor</u> plant <u>that has not been pre-selected for a particular phenotype</u>, which is then cloned into a cloning vector to form a genomic DNA library, with the proviso that the step does not include a preliminary selection step of the genomic DNA fragments;

2) introducing the genomic fragment from each of the genomic clones constituting the genomic DNA library separately and randomly into a <u>recipient plant that has not been pre-</u> <u>selected for a particular phenotype</u> to produce transgenic plants;

 cultivating the transgenic plants plant or progeny thereof to select a plant exhibiting an agriculturally advantageous phenotypic variation;

4) selecting the genomic DNA fragment, which was introduced in step (2) into the plant selected in step (3), as a purposed genomic DNA fragment; and

5) eptionally isolating and reintroducing the genomic DNA fragment selected in step (4) or a part thereof into another plant of any species to produce a plant to repeat steps (3) and (4), and selecting a genomic DNA fragment which produces a plant exhibiting an agriculturally advantageous phenotypic variation as a purposed genomic DNA fragment in each repetition; and

 cultivating the plant of step 5) to select a plant exhibiting said agriculturally advantageous phenotypic variation.

2. - 12. (Canceled)

Application No.: 10/576,693 Docket No.; 0230-0238PU\$1

Amendment dated September 15, 2010 Reply to Office Action of March 16, 2010

13. (Withdrawn) A method for producing a genomic DNA fragment capable of bringing

about an agriculturally advantageous phenotypic variation in plants comprising the steps of:

culturing E. coli cells containing a cloning vector carrying a genomic DNA fragment selected

by the method according to claim 1; and preparing the cloning vector amplified in the E. coli

cells along with the genomic DNA fragment.

14. (Withdrawn) A method for producing a genomic DNA fragment wherein the genomic

DNA fragment selected by the method according to claim 1 is used as a template and the

amplification of the fragment is conducted by a biochemical amplification method.

15. (Withdrawn) A method for producing a DNA fragment wherein the genomic DNA

fragment obtained by the method of claims 13 or 14 is digested with restriction enzyme(s),

16. (Canceled)

17. (Withdrawn and Currently Amended) A method for producing a plant having an

agriculturally advantageous phenotypic variation comprising the step of introducing a genomic

DNA fragment capable of bringing about an agriculturally advantageous phenotypic variation

in plants, wherein the genomic DNA fragment is screened by the method according to claim

1 and the method further produced by a method comprising comprises the steps of: culturing

E. coli cells containing a cloning vector carrying the genomic DNA fragment, and preparing

the cloning vector amplified in the E. coli cells along with the genomic DNA fragment.

3

Reply to Office Action of March 16, 2010

18. (Withdrawn) A method for producing a plant having an agriculturally advantageous

phenotypic variation according to claim 17 wherein the step of introducing a genomic DNA

fragment capable of bringing about an agriculturally advantageous phenotypic variation in

plants comprises the steps of: introducing the genomic fragment into a plant cell or tissue;

regenerating a complete plant from the plant cell; and cultivating the regenerated plant.

19. (Withdrawn) A method for producing a plant according to claim 18 wherein the

introduction of the genomic DNA fragment into a plant cell or tissue is conducted by a method

selected from the group consisting of biological introduction methods, physical introduction

methods and chemical introduction methods.

20. (Withdrawn) A method for producing a plant according to any one of claims 17 to 19,

wherein the genomic DNA fragment capable of bringing about an agriculturally advantageous

phenotypic variation is introduced in a plant of the same species as that of the plant from

which the genomic DNA fragment was derived.

21. (Withdrawn) A method for producing a plant according to any one of claims 17 to 19,

wherein genomic DNA fragment capable of introducing an agriculturally beneficial phenotypic

variation is introduced in a plant of a different species from the plant from which the genomic

DNA fragment was derived.

22. (Canceled)

23. (Withdrawn) A method for analyzing a genomic DNA fragment capable of bringing

4

GMM/SWG/eaw

Reply to Office Action of March 16, 2010

about an agriculturally advantageous phenotypic variation comprising the steps of: culturing

E. coli cells containing a cloning vector carrying a genomic DNA fragment selected by the

method according to claim 1; and preparing the cloning vectors amplified in the ${\rm E.}\ coli$ cells

along with the genomic DNA fragment, and reading the nucleotide sequence of the plant

genomic DNA fragment in the cloning vector.

24. (Withdrawn) A method for analyzing a DNA fragment comprising the step of

restricting the genomic DNA fragment selected according to claim 1.

25. (Withdrawn) A method for analyzing a DNA fragment wherein the genomic DNA

fragment selected by the method according to claim 1 is used as a template and the

amplification is conducted by a biochemical amplification method.

26. (Withdrawn) A method according to claim 24 or 25 wherein the analysis comprises the

step of reading the nucleotide sequence of the restriction product of the genomic DNA fragment

or the biochemically amplified product.

27. - 31. (Canceled)

32. (Currently Amended) The method according to claim 1or 31, wherein the size of the

selected genomic DNA fragment is 1 kb or greater provided that the DNA fragment can be

introduced into the cloning vector.

33. (Currently Amended) The method for screening according to elaim 1 or 31 claim 1,

5

GMM/SWG/eaw

Reply to Office Action of March 16, 2010

wherein step (2) comprises the sub-steps of: introducing the genomic fragment into the genome of a cell or tissue of the plant; regenerating a complete plant from the plant cell; and cultivating the regenerated plant.

- 34. (Previously Presented) The screening method according to claim 33, wherein the introduction of the genomic DNA fragment into a plant cell or tissue is conducted by a method selected from the group consisting of biological introduction methods, physical introduction methods and chemical introduction methods.
- 35. (Currently Amended) The screening method according to elaim-1-or-3+claim 1, wherein the agriculturally advantageous phenotypic mutation-variation in a plant gives rise to an increase or decrease of the size or the weight of at least a part of the plant or of at least a constituent thereof, an increase of growth rate or an excellent resistance against diseases or pests, under normal cultivation conditions, as compared with a case where the plant does not have the phenotypic variation.
- 36. (Currently Amended) The screening method according to elaim 1 or 31 claim 1, wherein the agriculturally advantageous phenotypic variation in a plant gives rise to an increase or decrease of the size or the weight of at least a part of the plant or of at least a constituent thereof, an increase of growth rate or an excellent resistance against diseases or pests, under conditions which are more stressful for the plant than normal conditions, as compared with a case where the plant does not have the phenotypic variation.
- 37. (Previously Presented) The screening method according to claim 35, wherein the plant

Application No.: 10/576,693 Amendment dated September 15, 2010

Reply to Office Action of March 16, 2010

transformed in step (2) is of the same species as that of the plant which supplied the genomic

DNA in step (1).

38. (Previously Presented) The screening method according to claim 36, wherein the plant

transformed in step (2) is of the same species from that of the plant which supplied the genomic

DNA in step (1).

39. (Previously Presented) The screening method according to claim 35, wherein the plant

transformed in step (2) is of a different species from that of the plant which supplied the genomic

DNA in step (1).

40. (Previously Presented) The screening method according to claim 36, wherein the plant

transformed in step (2) is of a different species from that of the plant which supplied the genomic

DNA in step (1).

41. (Currently Amended) The screening method according to claim 37, wherein the

optional-reintroduction of the genomic DNA fragment in step (5) is made into a plant of the same

species as that of the plant which was transformed in step (2).

42. (Currently Amended) The screening method according to claim 38, wherein the

optional-reintroduction of the genomic DNA fragment in step (5) is made into a plant of the same

species as that of the plant which was transformed in step (2).

43. (Currently Amended) The screening method according to clam 39, wherein the optional

7

GMM/SWG/eaw

Docket No.: 0230-0238PUS1

Reply to Office Action of March 16, 2010

reintroduction of the genomic DNA fragment in step (5) is made into a plant of the same species

as that of the plant which was transformed in step (2).

44. (Currently Amended) The screening method according to claim 40, wherein the

optional-reintroduction of the genomic DNA fragment in step (5) is made into a plant of the same

sphonaric introduction of the genomic DNA fragment in step (3) is made into a plant of the same

species as that of the plant which was transformed in step (2).

45. (Currently Amended) The screening method according to claim 37, wherein the

optional-reintroduction of the genomic DNA fragment on step (5) is made into a plant of a

different species from that of the plant which was transformed in step (2).

46. (Currently Amended) The screening method according to claim 38, wherein the

optional-reintroduction of the genomic DNA fragment in step (5) is made into a plant of a

different species from that of the plant which was transformed in step (2).

47. (Currently Amended) The screening method according to claim 39, wherein the

optional-reintroduction of the genomic DNA fragment in step (5) is made into a plant of a

different species from that of the plant which was transformed in step (2).

48. (Currently Amended) The screening method according to claim 40, wherein the

optional-reintroduction of the genomic DNA fragment in step (5) is made into a plant of a

different species from that of the plant which was transformed in step (2).

49. (NEW) A method for screening genomic DNA fragments capable of providing a plant

8

Reply to Office Action of March 16, 2010

with an agriculturally advantageous phenotypic variation, comprising the steps of:

preparing genomic DNA from a donor plant that has not been pre-selected for a
particular phenotype, which is then cloned into a cloning vector to form a genomic DNA library,
with the proviso that the step does not include a preliminary selection step of the genomic DNA

fragments;

2) introducing the genomic fragment from each of the genomic clones constituting the genomic DNA library separately and randomly into a recipient plant that has not been preselected for a particular phenotype, which is the same species as the donor plant, to produce a

transgenic plant;

 cultivating the transgenic plants or progeny thereof to select a plant exhibiting an agriculturally advantageous phenotypic variation;

4) selecting the genomic DNA fragment, which was introduced in step (2) into the plant

selected in step (3), as a purposed genomic DNA fragment; and

5) optionally introducing the genomic DNA fragment selected in step (4), or a part thereof, into another plant to repeat steps (3) and (4), and selecting a genomic DNA fragment which produces a plant exhibiting an agriculturally advantageous phenotypic variation as a

purposed genomic DNA fragment in each repetition.